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NOTICE OF ALLOWANCE AND FEE(S) DUE

30565

7590

12/14/2009

WOODARD, EMHARDT, MORIARTY, MCNETT & HENRY LLP 111 MONUMENT CIRCLE, SUITE 3700 INDIANAPOLIS, IN 46204-5137 EXAMINER

GUPTA, ANISH

ART UNIT PAPER NUMBER

1654 DATE MAILED: 12/14/2009

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/551,840	01/20/2006	Fabrizio Samaritani	7541-6	4228

TITLE OF INVENTION: LIQUID PHARMACEUTICAL FORMULATIONS OF FSH AND LH TOGETHER WITH A NON-IONIC SURFACTANT

L	APPLN. TYPE	SMALL ENTITY	ISSUE FEE DUE	PUBLICATION FEE DUE	PREV. PAID ISSUE FEE	TOTAL FEE(S) DUE	DATE DUE
	nonprovisional	NO	\$1510	\$300	\$0	\$1810	03/15/2010

THE APPLICATION IDENTIFIED ABOVE HAS BEEN EXAMINED AND IS ALLOWED FOR ISSUANCE AS A PATENT. PROSECUTION ON THE MERITS IS CLOSED. THIS NOTICE OF ALLOWANCE IS NOT A GRANT OF PATENT RIGHTS. THIS APPLICATION IS SUBJECT TO WITHDRAWAL FROM ISSUE AT THE INITIATIVE OF THE OFFICE OR UPON PETITION BY THE APPLICANT. SEE 37 CFR 1.313 AND MPEP 1308.

THE ISSUE FEE AND PUBLICATION FEE (IF REQUIRED) MUST BE PAID WITHIN THREE MONTHS FROM THE MAILING DATE OF THIS NOTICE OR THIS APPLICATION SHALL BE REGARDED AS ABANDONED. THIS STATUTORY PERIOD CANNOT BE EXTENDED. SEE 35 U.S.C. 151. THE ISSUE FEE DUE INDICATED ABOVE DOES NOT REFLECT A CREDIT FOR ANY PREVIOUSLY PAID ISSUE FEE IN THIS APPLICATION. IF AN ISSUE FEE HAS PREVIOUSLY BEEN PAID IN THIS APPLICATION (AS SHOWN ABOVE), THE RETURN OF PART B OF THIS FORM WILL BE CONSIDERED A REQUEST TO REAPPLY THE PREVIOUSLY PAID ISSUE FEE TOWARD THE ISSUE FEE NOW DUE.

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If the SMALL ENTITY is shown as YES, verify your current SMALL ENTITY status:

A. If the status is the same, pay the TOTAL FEE(S) DUE shown above.

B. If the status above is to be removed, check box 5b on Part B - Fee(s) Transmittal and pay the PUBLICATION FEE (if required) and twice the amount of the ISSUE FEE shown above, or

If the SMALL ENTITY is shown as NO:

A. Pay TOTAL FEE(S) DUE shown above, or

B. If applicant claimed SMALL ENTITY status before, or is now claiming SMALL ENTITY status, check box 5a on Part B - Fee(s) Transmittal and pay the PUBLICATION FEE (if required) and 1/2 the ISSUE FEE shown above.

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III. All communications regarding this application must give the application number. Please direct all communications prior to issuance to Mail Stop ISSUE FEE unless advised to the contrary.

IMPORTANT REMINDER: Utility patents issuing on applications filed on or after Dec. 12, 1980 may require payment of maintenance fees. It is patentee's responsibility to ensure timely payment of maintenance fees when due.

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Complete and send this form, together with applicable fee(s), to: Mail Mail Stop ISSUE FEE

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Note: A certificate of mailing can only be used for domestic mailings of the CURRENT CORRESPONDENCE ADDRESS (Note: Use Block 1 for any change of address) Fee(s) Transmittal. This certificate cannot be used for any other accompanying papers. Each additional paper, such as an assignment or formal drawing, must have its own certificate of mailing or transmission. 30565 7590 12/14/2009 Certificate of Mailing or Transmission WOODARD, EMHARDT, MORIARTY, MCNETT & HENRY LLIPhereby certify that this Fee(s) Transmittal is being deposited with the United States Postal Service with sufficient postage for first class mail in an envelope addressed to the Mail Stop ISSUE FEE address above, or being facsimile transmitted to the USPTO (571) 273-2885, on the date indicated below. 111 MONUMENT CIRCLE, SUITE 3700 INDIANAPOLIS, IN 46204-5137 (Depositor's name (Signature (Date APPLICATION NO. FILING DATE FIRST NAMED INVENTOR ATTORNEY DOCKET NO. CONFIRMATION NO. 10/551,840 01/20/2006 Fabrizio Samaritani 4228 TITLE OF INVENTION: LIQUID PHARMACEUTICAL FORMULATIONS OF FSH AND LH TOGETHER WITH A NON-IONIC SURFACTANT APPLN. TYPE SMALL ENTITY ISSUE FEE DUE PUBLICATION FEE DUE PREV. PAID ISSUE FEE TOTAL FEE(S) DUE DATE DUE nonprovisional NO \$1510 \$300 \$0 \$1810 03/15/2010 **EXAMINER** ART UNIT CLASS-SUBCLASS GUPTA, ANISH 1654 514-002000 1. Change of correspondence address or indication of "Fee Address" (37 CFR 1.363). 2. For printing on the patent front page, list (1) the names of up to 3 registered patent attorneys or agents OR, alternatively, ☐ Change of correspondence address (or Change of Correspondence Address form PTO/SB/122) attached. (2) the name of a single firm (having as a member a ☐ "Fee Address" indication (or "Fee Address" Indication form PTO/SB/47; Rev 03-02 or more recent) attached. Use of a Customer Number is required. registered attorney or agent) and the names of up to 2 registered patent attorneys or agents. If no name is listed, no name will be printed. 3. ASSIGNEE NAME AND RESIDENCE DATA TO BE PRINTED ON THE PATENT (print or type) PLEASE NOTE: Unless an assignee is identified below, no assignee data will appear on the patent. If an assignee is identified below, the document has been filed for recordation as set forth in 37 CFR 3.11. Completion of this form is NOT a substitute for filing an assignment. (A) NAME OF ASSIGNEE (B) RESIDENCE: (CITY and STATE OR COUNTRY) 4b. Payment of Fee(s): (Please first reapply any previously paid issue fee shown above) 4a. The following fee(s) are submitted: lssue Fee A check is enclosed. Publication Fee (No small entity discount permitted) Payment by credit card. Form PTO-2038 is attached. The Director is hereby authorized to charge the required fee(s), any deficiency, or credit any overpayment, to Deposit Account Number ______ (enclose an extra copy of this fo Advance Order - # of Copies _ (enclose an extra copy of this form). 5. Change in Entity Status (from status indicated above) ☐ b. Applicant is no longer claiming SMALL ENTITY status. See 37 CFR 1.27(g)(2). a. Applicant claims SMALL ENTITY status. See 37 CFR 1.27. NOTE: The Issue Fee and Publication Fee (if required) will not be accepted from anyone other than the applicant; a registered attorney or agent; or the assignee or other party in interest as shown by the records of the United States Patent and Trademark Office. Authorized Signature Date Typed or printed name Registration No. This collection of information is required by 37 CFR 1.311. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, Virginia 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, Virginia 22313-1450.

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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/551,840	01/20/2006	Fabrizio Samaritani	7541-6	4228
30565 75	30565 7590 12/14/2009		EXAMINER	
WOODARD, EM	MHARDT, MORIAR	GUPTA, ANISH		
	CIRCLE, SUITE 370	ART UNIT	PAPER NUMBER	
INDIANAPOLIS,	IN 46204-5137		1654	
			DATE MAILED: 12/14/200	9

Determination of Patent Term Adjustment under 35 U.S.C. 154 (b)

(application filed on or after May 29, 2000)

The Patent Term Adjustment to date is 0 day(s). If the issue fee is paid on the date that is three months after the mailing date of this notice and the patent issues on the Tuesday before the date that is 28 weeks (six and a half months) after the mailing date of this notice, the Patent Term Adjustment will be 0 day(s).

If a Continued Prosecution Application (CPA) was filed in the above-identified application, the filing date that determines Patent Term Adjustment is the filing date of the most recent CPA.

Applicant will be able to obtain more detailed information by accessing the Patent Application Information Retrieval (PAIR) WEB site (http://pair.uspto.gov).

Any questions regarding the Patent Term Extension or Adjustment determination should be directed to the Office of Patent Legal Administration at (571)-272-7702. Questions relating to issue and publication fee payments should be directed to the Customer Service Center of the Office of Patent Publication at 1-(888)-786-0101 or (571)-272-4200.

	Application No.	Applicant(s)	
	10/551,840	SAMARITANI ET AL.	
Notice of Allowability	Examiner	Art Unit	
	ANIIOLI OLIDTA	4054	
	ANISH GUPTA	1654	
The MAILING DATE of this communication apperature All claims being allowable, PROSECUTION ON THE MERITS IS herewith (or previously mailed), a Notice of Allowance (PTOL-85) NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT RIGHT of the Office or upon petition by the applicant. See 37 CFR 1.313	(OR REMAINS) CLOSED or other appropriate com IGHTS. This application is	o in this application. If not included munication will be mailed in due course. THIS	/e
1. This communication is responsive to <u>7-06-09</u> .			
2. \square The allowed claim(s) is/are $\underline{46-49,51-53,57-63,72-80,82-89}$	9,189-193,195-201 and 20	<u>03-217</u> .	
 3. Acknowledgment is made of a claim for foreign priority unally All b) Some* c) None of the: 1. Certified copies of the priority documents have 2. Certified copies of the priority documents have 	e been received.	, ,,	
3. Copies of the certified copies of the priority do	cuments have been recei	ved in this national stage application from the	
International Bureau (PCT Rule 17.2(a)).			
* Certified copies not received:			
Applicant has THREE MONTHS FROM THE "MAILING DATE" noted below. Failure to timely comply will result in ABANDONN THIS THREE-MONTH PERIOD IS NOT EXTENDABLE.		file a reply complying with the requirements	
4. A SUBSTITUTE OATH OR DECLARATION must be subm INFORMAL PATENT APPLICATION (PTO-152) which give			
5. CORRECTED DRAWINGS (as "replacement sheets") mus	st be submitted.		
(a) \square including changes required by the Notice of Draftspers	son's Patent Drawing Rev	iew (PTO-948) attached	
1) 🔲 hereto or 2) 🔲 to Paper No./Mail Date	•		
(b) ☐ including changes required by the attached Examiner's Paper No./Mail Date	s Amendment / Comment	or in the Office action of	
Identifying indicia such as the application number (see 37 CFR 1 each sheet. Replacement sheet(s) should be labeled as such in t			
6. DEPOSIT OF and/or INFORMATION about the depo attached Examiner's comment regarding REQUIREMENT			
Attachment(s)	5 Notice of	Informal Detant Application	
 Notice of References Cited (PTO-892) Notice of Draftperson's Patent Drawing Review (PTO-948) 		Informal Patent Application Summary (PTO-413),	
,	Paper N	o./Mail Date	
3. Information Disclosure Statements (PTO/SB/08), Paper No./Mail Date	7. 🛛 Examine	's Amendment/Comment	
4. Examiner's Comment Regarding Requirement for Deposit of Biological Material	8. 🛛 Examine	's Statement of Reasons for Allowance	
	9.	<u> </u>	
/Anish Gupta/			
Primary Examiner, Art Unit 1654			

Application/Control Number: 10/551,840

Art Unit: 1654

DETAILED ACTION

Page 2

EXAMINER'S AMENDMENT

1. An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with Thomas Henry on September 24, 2009.

The application has been amended as follows:

The following claims have been amended as follows:

46. A liquid pharmaceutical composition, comprising:

a follicle stimulating hormone or a variant thereof,

m-cresol,

a diluent, and

poloxamer 188,

the composition having a stability sufficient to avoid precipitation.

- 72. A liquid pharmaceutical composition, comprising:
 - a follicle stimulating hormone or a variant thereof,
 - a luteinising hormone or a variant thereof,
 - a bacteriostatic agent selected form the group consisting of phenol and m-cresol,

poloxamer 188, and

a diluent

Application/Control Number: 10/551,840

Art Unit: 1654

Page 3

wherein <u>either</u> the follicle stimulating hormone is human follicle stimulating hormone, <u>or</u> the luteinising hormone is human luteinising hormone, or <u>both</u> the follicle stimulating hormone is human follicle stimulating hormone and the luteinising hormone is human luteinising

hormone,

the composition having a stability sufficient to avoid precipitation.

73. A liquid pharmaceutical composition, comprising:

a follicle stimulating hormone or a variant thereof,

a luteinising hormone or a variant thereof,

a bacteriostatic agent selected form the group consisting of phenol and m-cresol,

poloxamer 188, and

a diluent

wherein <u>either</u> the follicle stimulating hormone is urinary human follicle stimulating hormone, <u>or</u> the luteinising hormone is urinary human luteinising hormone, or <u>both</u> the follicle stimulating hormone is urinary human follicle stimulating hormone and the luteinising hormone is urinary human luteinising hormone,

the composition having a stability sufficient to avoid precipitation.

74. A liquid pharmaceutical composition, comprising:

a follicle stimulating hormone or a variant thereof,

a luteinising hormone or a variant thereof,

a bacteriostatic agent selected form the group consisting of phenol and m-cresol,

poloxamer 188, and

Application/Control Number: 10/551,840

Art Unit: 1654

a diluent

wherein <u>either</u> the follicle stimulating hormone is recombinant human follicle stimulating hormone, <u>or</u> the luteinising hormone is recombinant human luteinising hormone, or both the follicle stimulating hormone is recombinant human follicle stimulating hormone and the recombinant luteinising hormone is human luteinising hormone, the composition having a stability sufficient to avoid precipitation.

Page 4

198. A liquid pharmaceutical composition, comprising:

a follicle stimulating hormone or a variant thereof,

phenol,

a diluent, and

poloxamer 188,

the composition having a stability sufficient to avoid precipitation.

The following is an examiner's statement of reasons for allowance:

The claims are drawn to a liquid composition comprising follicle stimulating hormone and poloxamer 188 as two of the specific components.

Hoffman et al. teaches a formulation comprising follicle stimulating hormone. The reference states "Other additives, such as a pharmaceutically acceptable solubilizers like Tween 20 (polyoxyethylene (20) sorbitan monopalmitate), Tween 40 (polyoxyethylene (20) sorbitan monopalmitate), Tween 80 (polyoxyethylene (20) sorbitan monopalmitate), Pluronic F68 (polyoxyethylene polyoxypropylene block copolymers), and PEG (polyethylene glycol) or non-ionic

surfactants such as polysorbate 20 or 80 or poloxamer 184 or 188, Pluronic.RTM. polyls, other block co-polymers, and chelators such as EDTA and EGTA may optionally be added to the formulations or compositions to reduce aggregation." (see paragraph [0100]). Thus, reading the reference, one would expect both poloxamer and tween 20 to behave similarly.

However, as Applicants specification asserts formulations with FSH and mixtures of FSH and LH with a Pluronic F68 (BASF, Pluronic F68 is also known as Poloxamer 188) obtain a stable formulation that avoids the problem of precipitation in the presence of a bacteriostatic agent, such as m-cresol and phenol. Precipitation, resulting in the formation of turbid or milky solutions occurs when TWEEN 20 is used with m-cresol or phenol (see page 12 of the specification). This is unexpected since one would have expected to observe the same "turbid or milky solution" for poloxamer 188 based on tween 20. Based on Hoffman et al.'s disclosure one would predict Tween 20 and poloxamer 188 to behave the same way. Furthermore, Hoffman et al. does not teach nor suggest the use of Poloxamer 188 over Tween 20, PEG or other surfactant listed. Indeed, Hoffman does not provide a single example that uses either Tween 20 or Poloxamer. Thus, based on Applicants disclosure that formulations with FSH and mixtures of FSH and LH with a Pluronic F68 (BASF, Pluronic F68 is also known as Poloxamer 188) obtain a stable formulation that avoids the problem of precipitation in the presence of a bacteriostatic agent, such as m-cresol and phenol, and the fact that Hoffman does not provide any specific motivation to choose Poloxamer 188 from the list of additives, the claims of the instant application are both novel and unobvious.

Any comments considered necessary by applicant must be submitted no later than the payment of the issue fee and, to avoid processing delays, should preferably accompany the issue fee. Such submissions should be clearly labeled "Comments on Statement of Reasons for Allowance."

2. Any inquiry concerning this communication or earlier communications from the examiner

should be directed to ANISH GUPTA whose telephone number is (571)272-0965. The examiner

can normally be reached on 5/4/9.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor,

Page 6

Tsang Cecilia can be reached on 571-272-0562. The fax phone number for the organization where

this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent

Application Information Retrieval (PAIR) system. Status information for published applications

may be obtained from either Private PAIR or Public PAIR. Status information for unpublished

applications is available through Private PAIR only. For more information about the PAIR system,

see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system,

contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like

assistance from a USPTO Customer Service Representative or access to the automated information

system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Anish Gupta/

Primary Examiner, Art Unit 1654